

EC Design Examination Certificate

The Notified Body

MEDCERT Zertifizierungs- und Prüfungsgesellschaft für die Medizin GmbH
Pilatuspool 2 – 20355 Hamburg – Germany

herewith certifies that the design of the medical device(s)

BeGraft Coronary Stent Graft System
with specifications listed in the appendix

of the manufacturer

Bentley InnoMed GmbH
Lotzenäcker 3
72379 Hechingen
Germany

Fulfills the below mentioned requirements of the **Council Directive 93/42/EEC**:

Annex II, section 4

This certificate assumes that MEDCERT has to be informed about any changes of the approved design. Changes need further approval by MEDCERT.

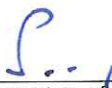
For the placing on the market of medical devices covered by this certificate, a separate EC certificate of conformity according to Annex II without section 4 of Council Directive 93/42/EEC is required.

Effective date: 2020-06-05

Expiry date: 2024-05-27

Report No.: PP13850FS02F
Process No.: PP – 13850
Certificate No.: 13850GB411200605

Hamburg, 2020-06-05



MEDCERT Certification Body
(Dr. Andreas Schich)

The certificate is only valid when provided entirely with all of its pages.
To verify the validity of this certificate, contact info@medcert.de.

MEDCERT Identification Number: 0482



Appendix of EC Design Examination Certificate

Process No.: PP – 13850

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Specifications

BeGraft Coronary Stent Graft System					
BG08250	BG12250	BG16250	BG18250	BG21250	BG24250
BG08275	BG12275	BG16275	BG18275	BG21275	BG24275
BG08300	BG12300	BG16300	BG18300	BG21300	BG24300
BG08350	BG12350	BG16350	BG18350	BG21350	BG24350
BG08400	BG12400	BG16400	BG18400	BG21400	BG24400
n.a.	n.a.	BG16450	BG18450	BG21450	BG24450
n.a.	n.a.	BG16500	BG18500	BG21500	BG24500

– End of list –

This appendix is integral part of the above-referenced certificate.
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